

# Raw Sequence Listing Error Summary

ERROR DETECTED	SUGGESTED CORRECTION	SERIAL NUMBER: <u>09/348,354</u>
ATTN: NEW RULES CASES: PLEASE DISREGARD ENGLISH "ALPHA" HEADERS, WHICH WERE INSERTED BY PTO SOFTWARE		
1 <input type="checkbox"/> Wrapped Nucleic	The number/text at the end of each line "wrapped" down to the next line. This may occur if your file was retrieved in a word processor after creating it. Please adjust your right margin to .3, as this will prevent "wrapping".	AUG 21 2000 TECH CENTER 1995/2000
2 <input type="checkbox"/> Wrapped Aminos	The amino acid number/text at the end of each line "wrapped" down to the next line. This may occur if your file was retrieved in a word processor after creating it. Please adjust your right margin to .3, as this will prevent "wrapping".	
3 <input type="checkbox"/> Incorrect Line Length	The rules require that a line not exceed 72 characters in length. This includes spaces.	
4 <input type="checkbox"/> Misaligned Amino Acid Numbering	The numbering under each 5th amino acid is misaligned. This may be caused by the use of tabs between the numbering. It is recommended to delete any tabs and use spacing between the numbers.	
5 <input type="checkbox"/> Non-ASCII	This file was not saved in ASCII (DOS) text, as required by the Sequence Rules. Please ensure your subsequent submission is saved in ASCII text so that it can be processed.	
6 <input type="checkbox"/> Variable Length	Sequence(s) <input type="checkbox"/> contain n's or Xaa's which represented more than one residue. As per the rules, each n or Xaa can only represent a single residue. Please present the maximum number of each residue having variable length and indicate in the (ix) feature section that some may be missing.	
7 <input type="checkbox"/> PatentIn ver. 2.0 "bug"	A "bug" in PatentIn version 2.0 has caused the <220>-<223> section to be missing from amino acid sequence(s) <input type="checkbox"/> . Normally, PatentIn would automatically generate this section from the previously coded nucleic acid sequence. Please manually copy the relevant <220>-<223> section to the subsequent amino acid sequence. This applies primarily to the mandatory <220>-<223> sections for Artificial or Unknown sequences.	
8 <input type="checkbox"/> Skipped Sequences (OLD RULES)	Sequence(s) <input type="checkbox"/> missing. If intentional, please use the following format for each skipped sequence: (2) INFORMATION FOR SEQ ID NO:X: (i) SEQUENCE CHARACTERISTICS:(Do not insert any headings under "SEQUENCE CHARACTERISTICS") (xi) SEQUENCE DESCRIPTION:SEQ ID NO:X: This sequence is intentionally skipped	
9 <input checked="" type="checkbox"/> Skipped Sequences (NEW RULES)	Please also adjust the "(iii) NUMBER OF SEQUENCES:" response to include the skipped sequence(s).  Sequence(s) <u>80</u> missing. If intentional, please use the following format for each skipped sequence. <210> sequence id number <400> sequence id number 000	
10 <input checked="" type="checkbox"/> Use of n's or Xaa's (NEW RULES)	Use of n's and/or Xaa's have been detected in the Sequence Listing. Use of <220> to <223> is MANDATORY if n's or Xaa's are present. In <220> to <223> section, please explain location of n or Xaa, and which residue n or Xaa represents.	
11 <input type="checkbox"/> Use of <213>Organism (NEW RULES)	Sequence(s) <input type="checkbox"/> are missing this mandatory field or its response.	
12 <input type="checkbox"/> Use of <220>Feature (NEW RULES)	Sequence(s) <input type="checkbox"/> are missing the <220>Feature and associated headings. Use of <220> to <223> is MANDATORY if <213>ORGANISM is "Artificial" or "Unknown" Please explain source of genetic material in <220> to <223> section. (See "Federal Register," 6/01/98, Vol. 63, No. 104, pp. 29631-32) (Sec. 1.823 of new Rules)	
13 <input type="checkbox"/> PatentIn ver. 2.0 "bug"	Please do not use "Copy to Disk" function of PatentIn version 2.0. This causes a corrupted file, resulting in missing mandatory numeric identifiers and responses (as indicated on raw sequence listing). Instead, please use "File Manager" or any other means to copy file to floppy disk.	

Does Not Comply  
Corrected Diskette Needed

1633

RAW SEQUENCE LISTING  
PATENT APPLICATION: US/09/348,354

DATE: 08/15/2000  
TIME: 15:21:47

Input Set : A:\PTO.txt  
Output Set: N:\CRF3\08152000\I348354.raw

3 <110> APPLICANT: Havenga, Menzo  
5 <120> TITLE OF INVENTION: Chimeric Adenoviruses  
7 <130> FILE REFERENCE: 2183-4123us  
9 <140> CURRENT APPLICATION NUMBER: 09/348,354  
10 <141> CURRENT FILING DATE: 1999-07-07  
E--> 12 <160> NUMBER OF SEQ ID NOS: 85  
14 <170> SOFTWARE: Patentin Ver. 2.1

## ERRORED SEQUENCES

3394 <210> SEQ ID NO: 79  
3395 <211> LENGTH 21 → 35  
3396 <212> TYPE: DNA  
3397 <213> ORGANISM: Primer/Oligonucleotide  
3399 <400> SEQUENCE: 79  
E--> 3400 gctcgatgtcaatgagggcg tgcgggtgggtc tcttc  
E--> 3403 <210> SEQ ID NO: 81

Sequence # 80 missing, causing  
miscount of sequences.

85 input, 84 counted.

Error in base count → 35  
14

VERIFICATION SUMMARY  
PATENT APPLICATION: US/09/348,354

DATE: 08/15/2000  
TIME: 15:21:48

Input Set : A:\PTO.txt  
Output Set: N:\CRF3\08152000\I348354.raw

L:298 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16  
L:298 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16  
L:298 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16  
L:298 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16  
L:298 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:16  
L:301 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16  
L:301 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16  
L:301 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16  
L:301 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16  
M:340 Repeated in SeqNo=16  
L:304 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16  
L:304 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16  
L:304 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16  
L:304 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16  
L:307 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16  
L:307 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16  
L:307 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16  
L:307 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16  
L:370 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16  
L:370 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16  
L:370 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16  
L:370 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16  
L:696 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:21  
L:696 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:21  
L:696 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:21  
L:696 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:21  
L:696 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:21  
L:976 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:976 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:976 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25  
L:976 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
L:976 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:25  
L:979 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:979 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:979 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25  
L:979 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
M:340 Repeated in SeqNo=25  
L:982 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:982 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:982 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25  
L:982 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
L:991 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:991 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:991 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25  
L:991 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
L:1000 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:1000 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:1000 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25

## VERIFICATION SUMMARY

PATENT APPLICATION: US/09/348,354

DATE: 08/15/2000

DATE: 05/15/20

Input Set : A:\PTO.txt  
Output Set: N:\CRF3\08152000\I348354.raw

L:1000 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
L:1006 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:25  
L:1006 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:25  
L:1006 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:25  
L:1006 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:25  
L:1532 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:32  
L:1532 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:32  
L:1532 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:32  
M:340 Repeated in SeqNo=32  
L:1723 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:34  
L:1760 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:35  
M:340 Repeated in SeqNo=35  
L:2252 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:41  
L:3400 M:254 E: No. of Bases conflict, LENGTH:Input:14 Counted:35 SEQ:79  
L:3400 M:252 E: No. of Seq. differs, <211>LENGTH:Input:21 Found:35 SEQ:79  
L:3403 M:214 E: (33) Seq.# missing, SEQ ID NO:80  
L:12 M:203 E: No. of Seq. differs, <160> Number Of Sequences:Input (85) Counted (84)

<210> 79  
<211> 21  
<212> DNA  
<213> Primer/Oligonucleotide

<400> 79  
gctcgatgt acaatgaggcg tgcgggtggtg tcttc  
14

→ Sequence # 80? If skipped intentionally,  
<210> 81 please use correct format as shown in  
<211> 34  
<212> DNA # 9 on Error Summary Sheet.  
<213> Primer/Oligonucleotide

<400> 81  
gctcgactta agttagaagg tgcgactgga aagc

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As per new sequence  
rules, all "n's" or "Xaa's" in  
sequences must have <220> to  
<223> features. See #

<210> 16  
<211> 391  
<212> PRT  
<213> Human Adenovirus 13 Fiber Protein

<400> 16

Xaa Xaa Xaa Xaa Xaa Ser Ala Pro Thr Ile Phe Met Leu Leu Gln Met  
1 5 10 15

Lys Arg Ala Arg Ser Ser Xaa Asp Thr Phe Asn Pro Val Tyr Pro Tyr  
20 25 30

Gly Tyr Ala Arg Asn Gln Asn Ile Xaa Phe Xaa Thr Pro Pro Phe Val  
35 40 45

Xaa Ser Asp Gly Phe Lys Asn Phe Pro Pro Gly Val Leu Ser Leu Lys  
50 55 60

Leu Ala Asp Pro Ile Thr Ile Ala Asn Gly Asp Val Ser Leu Lys Val  
65 70 75 80

Gly Gly Gly Leu Thr Leu Gln Glu Gly Ser Leu Thr Val Asp Pro Lys  
85 90 95

Ala Pro Leu Gln Leu Ala Asn Asp Lys Lys Leu Glu Leu Val Tyr Asp  
100 105 110

Asp Pro Phe Glu Val Ser Thr Asn Lys Leu Ser Leu Lys Val Gly His  
115 120 125

Gly Leu Lys Val Leu Asp Asp Lys Ser Ala Gly Gly Leu Lys Asp Leu  
130 135 140

Ile Gly Lys Leu Val Val Leu Thr Gly Lys Gly Ile Gly Ile Glu Asn  
145 150 155 160

Leu Gln Asn Asp Asp Gly Ser Ser Arg Gly Val Gly Ile Asn Val Arg  
165 170 175

Leu Gly Thr Asp Gly Gly Leu Ser Phe Asp Arg Lys Gly Glu Leu Val  
180 185 190

Ala Trp Asn Arg Lys Asp Asp Arg Arg Thr Leu Trp Thr Thr Pro Asp  
195 200 205

Pro Ser Pro Asn Cys Lys Ala Glu Thr Glu Lys Asp Ser Lys Leu Thr  
210 215 220

Leu Val Leu Thr Lys Cys Gly Ser Gln Ile Leu Ala Thr Val Ser Ile  
225 230 235 240

Ile Val Leu Lys Gly Lys Tyr Glu Phe Val Lys Lys Glu Thr Glu Pro  
245 250 255

Lys Ser Phe Asp Val Lys Leu Leu Phe Asp Ser Lys Gly Val Leu Leu  
260 265 270

Summary Sheet.

Please check  
entire sequence  
listing as this  
same error  
has been  
indicated  
throughout.

The types of errors shown exist throughout the Sequence Listing. Please check  
subsequent sequences for similar errors.